

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: June 25, 2003, 14:20:41 ; Search time 33 Seconds

(without alignments)
444.169 Million cell updates/sec

Title: US-09-622-613b-15

Perfect score: 602

Sequence: 1 QMNAFPOKHIIITPILICNT.....ICVKCENQYFVHAGIGRCP 110

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 10

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

1: /SID2/gcgdata/geneseq/emb1/AA1980.DAT:*
2: /SID2/gcgdata/geneseq/emb1/AA1981.DAT:*
3: /SID2/gcgdata/geneseq/emb1/AA1982.DAT:*
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19: /SID2/gcgdata/geneseq/emb1/AA1998.DAT:*
20: /SID2/gcgdata/geneseq/emb1/AA1999.DAT:*
21: /SID2/gcgdata/geneseq/emb1/AA2000.DAT:*
22: /SID2/gcgdata/geneseq/emb1/AA2001.DAT:*
23: /SID2/gcgdata/geneseq/emb1/AA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	602	100.0	110	20	AAV28872
2	602	100.0	111	20	AAV28873
3	597	99.2	110	20	AAV28877
4	597	99.2	111	20	AAV28878
5	596	99.0	110	20	AAV28874
6	596	99.0	111	20	AAV28876
7	588.5	97.8	111	20	AAV33321
8	284.5	47.3	104	18	AAW06544
9	281.5	46.8	104	20	AAV28865
10	281.5	46.8	105	20	AAV28867

11	281.5	46.8	127	20	AAV28879	Rana pipiens Clone
12	278.5	46.3	104	20	AAV28866	Recombinant RAPRL1
13	278.5	46.3	105	20	AAV28869	Recombinant Met(-1)
14	277.5	46.1	104	18	AAW30301	Recombinant onc pr
15	277.5	46.1	104	22	AAW31666	Amino acid sequenc
16	277.5	46.1	105	20	AAV39400	Recombinant frog O
17	277.5	46.1	104	18	AAW35126	R. pipiens recombi
18	276.5	45.9	104	20	AAV28870	Recombinant RAPRL1
19	276.5	45.9	105	20	AAV28871	Recombinant Met(-1)
20	274.5	45.6	104	12	AAW12344	Protein with activ
21	274.5	45.6	104	15	AAW47303	ONCONASE (pharmac
22	274.5	45.6	104	17	AAW0736	Protein derived fr
23	274.5	45.6	104	18	AAW06543	Antitumor protein
24	274.5	45.6	104	18	AAW14065	Onconase (RTM) pro
25	274.5	45.6	104	20	AAV33322	Frog onconase prot
26	274.5	45.6	104	20	AAW88233	Rana pipiens RNase
27	274.5	45.6	105	18	AAW35123	R. pipiens recombi
28	274.5	45.6	105	18	AAW35125	R. pipiens recombi
29	274.5	45.6	358	18	AAW35130	R. pipiens recombi
30	272.5	45.3	104	22	AAW31667	Amino acid sequenc
31	272.5	45.3	106	18	AAW35122	R. pipiens recombi
32	272.5	45.3	107	18	AAW35117	R. pipiens recombi
33	272.5	45.3	112	18	AAW35118	R. pipiens recombi
34	272.5	45.3	251	18	AAW35134	R. pipiens recombi
35	272.5	45.3	254	18	AAW35135	R. pipiens recombi
36	272.5	45.3	355	18	AAW35129	R. pipiens recombi
37	272.5	45.3	355	18	AAW35133	R. pipiens recombi
38	272.5	45.3	366	18	AAW35132	R. pipiens recombi
39	271.5	45.1	104	18	AAW30302	Recombinant onc pr
40	267.5	44.4	104	18	AAW18224	Antitumor generi
41	267.5	44.4	105	18	AAW35115	R. pipiens recombi
42	267.5	44.4	105	18	AAW35116	R. pipiens recombi
43	263.5	43.8	358	18	AAW35127	R. pipiens recombi
44	263.5	43.8	365	18	AAW35131	R. pipiens recombi
45	249.5	41.4	107	18	AAW35120	R. pipiens recombi

ALIGNMENTS

RESULT 1
AAV28872
ID AAV28872 standard; Protein; 110 AA.
XX
AC AAV28872;
XX
DT 25-JAN-2000 (first entry)
XX
XX Rana catesbeiana oocyte ribonuclease (RacOR1) amino acid sequence.
DE Rana catesbeiana oocyte ribonuclease: RacOR1; covalently bound; CD22;
KW LL2 antibody; ligand binding moiety; cancerous B cell; Kaposi's Sarcoma;
KW human chorionic gonadotropin; hCG; recombinant ribonuclease; bullfrog;
KW signal peptide; cytotoxic fusion protein; cancer; autoimmune disease;
KW RNase.
KW Rana catesbeiana.
XX
OS Synthetic.
XX
XX WO9950398-A2.
XX
XX PD 07-OCT-1999.
XX
XX PF 26-MAR-1999; 99WO-US06641.
XX
XX PR 27-MAR-1998; 98US-0079751.
XX
XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
XX PI Newton DL, Rybak SM;
XX
XX DR WPI: 1999-610847/52.
XX
XX DR N-PSDB; AA208130.

XX New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases
 XX
 PS Claim 22: Page 62; 71pp: English.
 XX
 CC The present sequence is a Rana catesbeiana oocyte ribonuclease (RacOR1)
 CC protein encoded by a cDNA modified for expression in *E. coli*. Carboxy
 CC terminal end of RacOR1 has a covalently bound ligand binding moiety,
 CC which can be a IL2 antibody directed against CD22 on cancerous B cells
 CC or human chorionic gonadotropin (hCG) effective against Kaposi's
 CC Sarcoma cells. Recombinant ribonucleases can be expressed in bacteria
 CC without an N-terminal methionine due to the presence of a signal peptide
 CC that is cleaved by bacteria. The soluble expression of ribonuclease
 CC allows the proteins to be fused in-frame with ligand binding moieties to
 CC form cytotoxic fusion proteins. They can be used for treatment of cancer
 CC and autoimmune diseases.
 XX
 SQ Sequence 110 AA:
 Query Match 100.0%; Score 602; DB 20; Length 110;
 Best Local Similarity 100.0%; Pred. No. 2.3e-61;
 Matches 110; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QNATFOCKHINPTICNTIMDNIIYVGQCKRVNFTFISSATYKAICTGVINNVL 60
 Db 1 QNATFOCKHINPTICNTIMDNIIYVGQCKRVNFTFISSATYKAICTGVINNVL 60
 QY 61 STTRFQNTCTRTSITPRCPYSSRTETNYICVGCENQYVHFAGIGRCP 110
 Db 61 STTRFQNTCTRTSITPRCPYSSRTETNYICVGCENQYVHFAGIGRCP 110
 RESULT 2
 ID AAY28873 standard; Protein: 111 AA.
 AC AAY28873:
 XX 25-JAN-2000 (first entry)
 DT
 DE Recombinant Met(-1) RacOR1.
 XX
 KW Recombinant Met(-1) Rana catesbeiana oocyte ribonuclease; RacOR1; CD22;
 KW covalently bound; IL2 antibody; ligand binding moiety; cancerous B cell;
 KW Kaposi's sarcoma; human chorionic gonadotropin; hCG; signal peptide;
 KW recombinant ribonuclease; cytotoxic fusion protein; cancer; bullfrog;
 KW Rnase; autoimmune disease.
 XX
 OS Rana catesbeiana.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 1 /note="Met not found in wild type RacOR1"
 FT
 XX WO9950398-A2.
 PN 07-OCT-1999.
 PD
 XX 26-MAR-1999; 99WO-US06641.
 PE
 XX 27-MAR-1998; 98US-0079751.
 PR
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX Newton DL, Rybak SM;
 XX WPI; 1999-610847/52.
 DR N-PSDB; AA208131.
 XX
 PT New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases

XX Claim 22: Page 63; 71pp: English.
 PS
 XX
 CC The present sequence is a recombinant Rana catesbeiana oocyte
 CC ribonuclease (RacOR1) protein with Met at position 1. Carboxy terminal
 CC end of recombinant RacOR1 has a covalently bound ligand binding moiety,
 CC which can be a IL2 antibody directed against CD22 on cancerous B cells or
 CC human chorionic gonadotropin (hCG) effective against Kaposi's sarcoma
 CC cells. Recombinant ribonucleases can be expressed in bacteria without an
 CC N-terminal methionine due to the presence of a signal peptide that is
 CC cleaved by bacteria. The soluble expression of ribonuclease allows the
 CC proteins to be fused in-frame with ligand binding moieties to form
 CC cytotoxic fusion proteins. They can be used for treatment of cancer and
 CC autoimmune diseases.
 XX
 SQ Sequence 111 AA:
 Query Match 100.0%; Score 602; DB 20; Length 111;
 Best Local Similarity 100.0%; Pred. No. 2.3e-61;
 Matches 110; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QNATFOCKHINPTICNTIMDNIIYVGQCKRVNFTFISSATYKAICTGVINNVL 60
 Db 2 QNATFOCKHINPTICNTIMDNIIYVGQCKRVNFTFISSATYKAICTGVINNVL 61
 QY 61 STTRFQNTCTRTSITPRCPYSSRTETNYICVGCENQYVHFAGIGRCP 110
 Db 62 STTRFQNTCTRTSITPRCPYSSRTETNYICVGCENQYVHFAGIGRCP 111
 RESULT 3
 ID AAY28877 standard; Protein: 110 AA.
 AC AAY28877:
 XX 25-JAN-2000 (first entry)
 DT
 DE Recombinant RacOR1 Glniser amino acid sequence.
 XX
 KW Recombinant Rana catesbeiana oocyte ribonuclease; RacOR1 Glniser; CD22;
 KW covalently bound; IL2 antibody; ligand binding moiety; cancerous B cell;
 KW bullfrog; Kaposi's sarcoma; human chorionic gonadotropin; hCG; Rnase;
 KW signal peptide; recombinant ribonuclease; cytotoxic fusion protein;
 KW cancer; autoimmune disease.
 XX
 OS Rana catesbeiana.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 1 /note="Wild type Gln replaced with Ser"
 FT
 XX WO9950398-A2.
 PN 07-OCT-1999.
 PD
 XX 26-MAR-1999; 99WO-US06641.
 PE
 XX 27-MAR-1998; 98US-0079751.
 PR
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX Newton DL, Rybak SM;
 XX WPI; 1999-610847/52.
 DR N-PSDB; AA208134.
 XX
 PT New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases
 PS Claim 22: Page 67; 71pp: English.
 XX

CC The present sequence is a recombinant Rana catesbeiana oocyte
CC ribonuclease (RacOR1) protein with Gln1Ser. Carboxy terminal end of
CC recombinant RacOR1 has a covalently bound ligand binding moiety, which
CC can be a LL2 antibody directed against CD22 on cancerous B cells or
CC human chorionic gonadotropin (hCG) effective against Kaposi's sarcoma
CC cells. Recombinant ribonucleases can be expressed in bacteria without an
CC N-terminal methionine due to the presence of a signal peptide that is
CC cleaved by bacteria. The soluble expression of ribonuclease allows the
CC proteins to be fused in-frame with ligand binding moieties to form
CC cytotoxic fusion proteins. They can be used for treatment of cancer and
CC autoimmune diseases.

XX Sequence 110 AA:

Query Match 99.2%; Score 597; DB 20; Length 110;
Best Local Similarity 100.0%; Pred. No. 8.7e-61;
Matches 109; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 NMATFOCKHILNPIICNTIMDNINIVYGCKRVNFIISATTVAKICTGVINMVL 61
DB 2 NMATFOCKHILNPIICNTIMDNINIVYGCKRVNFIISATTVAKICTGVINMVL 61
OY 62 TTRFQALNTCTRTSITPRCPYSSRTETNYICVKCENQYPVHFAIGRCP 110
DB 62 TTRFQALNTCTRTSITPRCPYSSRTETNYICVKCENQYPVHFAIGRCP 110

RESULT 4

AAI28878 standard; Protein: 111 AA.

XX AAY28878;

XX 25-JAN-2000 (first entry)

XX Recombinant Met(-1) RacOR1 Gln1Ser amino acid sequence.

XX Recombinant Met(-1) Rana catesbeiana oocyte ribonuclease Gln1Ser; RacOR1;
KW covalently bound; LL2 antibody; ligand binding moiety; cancerous B cell;
KW Kaposi's sarcoma; human chorionic gonadotropin; hCG; signal peptide;
KW recombinant ribonuclease; cytotoxic fusion protein; cancer; bullfrog;
KW CD22; RNase; autoimmune disease.

XX Rana catesbeiana.
OS Synthetic.

XX Key Location/Qualifiers

FT MISC-difference 1 /note= "Met not found in wild type RacOR1"

FT MISC-difference 2 /note= "Wild type Gln replaced with Ser"

XX W09950398-A2.

XX 07-OCT-1999.

XX 26-MAR-1999; 99WO-US06641.

XX 27-MAR-1998; 98US-0079751.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Newton DL, Rybak SM;

XX WPI: 1999-610847/52.

XX N-PSDB: AA208135.

XX New recombinant ribonucleases, used for killing target cells, e.g. for

XX treating cancers, viral infections or autoimmune diseases

XX Claim 22; Page 68; 71pp; English.

XX The present sequence is a recombinant Rana catesbeiana ribonuclease

CC (RacOR1) protein with Met at position 1 and Gln2Ser. Carboxy terminal end
CC of recombinant RacOR1 has a covalently bound ligand binding moiety, which
CC can be a LL2 antibody directed against CD22 on cancerous B cells or human
CC chorionic gonadotropin (hCG) effective against Kaposi's sarcoma cells.
CC Recombinant ribonucleases can be expressed in bacteria without an N-
CC terminal methionine due to the presence of a signal peptide that is
CC cleaved by bacteria. The soluble expression of ribonuclease allows the
CC proteins to be fused in-frame with ligand binding moieties to form
CC cytotoxic fusion proteins. They can be used for treatment of cancer and
CC autoimmune diseases.

XX Sequence 111 AA:

Query Match 99.2%; Score 597; DB 20; Length 111;
Best Local Similarity 100.0%; Pred. No. 8.8e-61;
Matches 109; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 NMATFOCKHILNPIICNTIMDNINIVYGCKRVNFIISATTVAKICTGVINMVL 61
DB 3 NMATFOCKHILNPIICNTIMDNINIVYGCKRVNFIISATTVAKICTGVINMVL 62
OY 62 TTRFQALNTCTRTSITPRCPYSSRTETNYICVKCENQYPVHFAIGRCP 110
DB 63 TTRFQALNTCTRTSITPRCPYSSRTETNYICVKCENQYPVHFAIGRCP 111

RESULT 5

AAI28874 standard; Protein: 110 AA.

XX AAY28874;

XX 25-JAN-2000 (first entry)

XX Recombinant RacOR1 Met22Leu Met57Leu amino acid sequence.

XX Recombinant Rana catesbeiana oocyte ribonuclease; covalently bound;
KW RacOR1 Met22Leu Met57Leu; LL2 antibody; ligand binding moiety; CD22;
KW cancerous B cell; Kaposi's sarcoma; human chorionic gonadotropin; hCG;
KW signal peptide; recombinant ribonuclease; cytotoxic fusion protein;
KW cancer; bullfrog; RNase; autoimmune disease.

XX Rana catesbeiana.
OS Synthetic.

XX Key Location/Qualifiers

FT MISC-difference 22 /note= "Wild type Met replaced with Leu"

FT MISC-difference 57 /note= "Wild type Met replaced with Leu"

XX W09950398-A2.

XX 07-OCT-1999.

XX 26-MAR-1999; 99WO-US06641.

XX 27-MAR-1998; 98US-0079751.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Newton DL, Rybak SM;

XX WPI: 1999-610847/52.

XX N-PSDB: AA208132.

XX New recombinant ribonucleases, used for killing target cells, e.g. for

XX treating cancers, viral infections or autoimmune diseases

XX Claim 22; Page 64; 71pp; English.

XX The present sequence is a recombinant Rana catesbeiana oocyte

XX ribonuclease (RacOR1) protein with Met22Leu Met57Leu. Carboxy terminal

CC end of recombinant RacOR1 has a covalently bound ligand binding moiety,
 CC which can be a LL2 antibody directed against CD22 on cancerous B cells,
 CC or human chorionic gonadotrophin (hCG) effective against Kaposi's sarcoma
 CC cells. Recombinant ribonucleases can be expressed in bacteria without an
 CC N-terminal methionine due to the presence of a signal peptide that is
 CC cleaved by bacteria. The soluble expression of ribonuclease allows the
 CC proteins to be fused in-frame with ligand binding moieties to form
 CC cytotoxic fusion proteins. They can be used for treatment of cancer and
 CC autoimmune diseases.

XX Sequence 110 AA:
 SQ

Query Match 99.0%; Score 596; DB 20; Length 110;
 Best Local Similarity 98.2%; Pred. No. 1.1e-60;
 Matches 108; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 QNMATFOOKHIINTPLICNTIMDNNTIYVGGCKRVNFTIISATVKAICTGVINLV 60
 DB 1 QNMATFOOKHIINTPLICNTIMDNNTIYVGGCKRVNFTIISATVKAICTGVINLV 60

QY 61 STTRFQNLNCTRTSITPRCPYSSRTETNYICVKCENQVPHFAGIGRCP 110
 DB 61 STTRFQNLNCTRTSITPRCPYSSRTETNYICVKCENQVPHFAGIGRCP 110

RESULT 6
 AAY28876
 ID AAY28876 standard; Protein: 111 AA.
 AC AAY28876;
 XX
 XX 25-JAN-2000 (first entry)
 DT
 XX
 DE Recombinant Met(-1) RacOR1 Met22Leu Met57Leu-(His)6 protein.
 XX
 XX Met(-1) Rana catesbeiana ribonuclease Met22Leu Met57Leu-(His)6; RacOR1;
 KW recombinant; CD22; covalently bound; LL2 antibody; ligand binding moiety;
 KW cancerous B cell; Kaposi's sarcoma; human chorionic gonadotrophin; hCG;
 KW signal peptide; recombinant ribonuclease; cytotoxic fusion protein;
 KW cancer; bullfrog; RNase; autoimmune disease.
 XX
 XX Rana catesbeiana.
 OS Synthetic.
 OS
 XX
 FH Key Location/Qualifiers
 FT MISC-difference 1 /note= "(His)6 histidine tag attached to N-terminal Met"
 FT MISC-difference 1 /note= "Met not found in wild type RacOR1"
 FT MISC-difference 23 /note= "Wild type Met replaced with Leu"
 FT MISC-difference 58 /note= "Wild type Met replaced with Leu"
 FT
 XX
 XX W09950398-A2.
 PN
 XX
 XX 07-OCT-1999.
 PD
 XX
 XX 26-MAR-1999; 99WO-US06641.
 PF
 XX
 XX 27-MAR-1998; 98US-0079751.
 PR
 XX
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX
 XX Newton DL, Rybak SM;
 PI
 XX
 XX WPI: 1999-610847/52.
 DR
 XX
 XX N-PSDB: AA208133.
 DR
 XX
 XX New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases -
 PT
 XX
 PS Claim 22; Page 66; 71pp; English.

XX
 CC The present sequence is a recombinant Rana catesbeiana oocyte
 CC ribonuclease (RacOR1) protein with Met at position 1 attached to a
 CC (His)6 tag, Met23Leu and Met58Leu. Carboxy terminal end of recombinant
 CC RacOR1 has a covalently bound ligand binding moiety, which can be a LL2
 CC antibody directed against CD22 on cancerous B cells or human chorionic
 CC gonadotrophin (hCG) effective against Kaposi's sarcoma cells. Recombinant
 CC ribonucleases can be expressed in bacteria without an N-terminal
 CC methionine due to the presence of a signal peptide that is cleaved by
 CC bacteria. The soluble expression of ribonuclease allows the proteins to
 CC be fused in-frame with ligand binding moieties to form cytotoxic fusion
 CC proteins. They can be used for treatment of cancer and autoimmune
 CC diseases.

XX Sequence 111 AA:
 SQ

Query Match 99.0%; Score 596; DB 20; Length 111;
 Best Local Similarity 98.2%; Pred. No. 1.1e-60;
 Matches 108; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 QNMATFOOKHIINTPLICNTIMDNNTIYVGGCKRVNFTIISATVKAICTGVINLV 60
 DB 2 QNMATFOOKHIINTPLICNTIMDNNTIYVGGCKRVNFTIISATVKAICTGVINLV 61

QY 61 STTRFQNLNCTRTSITPRCPYSSRTETNYICVKCENQVPHFAGIGRCP 110
 DB 62 STTRFQNLNCTRTSITPRCPYSSRTETNYICVKCENQVPHFAGIGRCP 111

RESULT 7
 AAY33321
 ID AAY33321 standard; Protein: 111 AA.
 AC AAY33321;
 XX
 XX 29-NOV-1999 (first entry)
 DT
 XX
 DE Frog lectin protein fragment.
 XX
 XX Cytotoxic; RNase; ribonuclease; pancreatic; antibody; light chain;
 KW heavy chain; cell surface marker; treatment; tumor; viral infection;
 KW parasite infection; immune dysfunctional cell; autoimmune disease;
 KW contraceptive; cell separation; transplantation; bone marrow ablation;
 KW leukemia cell; T-cell; graft-versus-host disease; bullfrog; lectin.
 XX
 XX Rana catesbeiana.
 OS
 OS
 XX
 PN US9595073-A.
 PN
 XX
 XX 21-SEP-1999.
 PD
 XX
 XX 09-JUL-1997; 97US-0891848.
 PF
 XX
 XX 22-SEP-1993; 93US-0125462.
 PR
 XX 22-OCT-1991; 91US-0779195.
 PR 20-APR-1990; 90US-0510696.
 PR 04-FEB-1993; 93US-0014082.
 PR
 XX
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX
 XX Rybak SM, Newton DL, Nicholls PJ, Youle RJ;
 PI
 XX
 XX WPI: 1999-560488/47.
 DR
 XX
 XX Recombinantly fused pancreatic RNase-targeting proteins useful for
 PT treating tumors, infections, immune or autoimmune disorders and as a
 PT contraceptive -
 PT
 XX
 XX Example 3; Fig 19; 47pp; English.
 PS
 XX
 XX This invention describes a novel nucleic acid construct comprising
 CC sequences encoding functional pancreatic RNase and a second protein
 CC (preferably the light and heavy chains of an antibody) which binds a

CC other 1 AAG6543) that have been isolated from Rana pliens oocytes.
CC Both proteins have a blocked amino terminal group and are essentially
CC free of carbohydrates. The proteins are used to treat tumours. Use of
CC the peptides has fewer disadvantages than chemotherapy, radiotherapy
CC and surgery in the treatment of tumours.

	Matches	55: conservative	15: mismatches	32: indels	9: gaps
Qy	1	ONMATEPQKHILIN-PLICNTIMDNINLYIGGCGKRVNFFITISATPTKALCTGVI-MNN	58		
Db	1	QDMLFEKKHILITNTRDDCNINISTNLF---HCKRKNFFITSRPEPAVALKGIILASN	56		

OY 59 VLSSTRQALNTCTRTSTTPRCPPYSSRTETNYICVGCENQYPVHFGIGRC 109
 DB 57 VLTTSERYLSDC---NWTSRPCKYKLLKSTNTFCVTCENQAPVHFGVGHG 104

RESULT 10

AAy28867
 ID AAY28867 standard; Protein: 105 AA.

AC AAY28867;
 DT 25-JAN-2000 (first entry)

DE Recombinant Met(-1) RapLRI.
 KW Recombinant Met(-1) Rana pipiens ribonuclease; RapLRI; CD22; RNase;
 KM covalently bound; IL2 antibody; ligand binding moiety; cancerous B cell;
 KM Kaposi's sarcoma; human chorionic gonadotrophin; hCG; signal peptide;
 KM recombinant ribonuclease; cytotoxic fusion protein; cancer; frog;
 KM autoimmune disease.

OS Rana pipiens.
 OS Synthetic.

FT Key Location/Qualifiers
 FT Misc-difference 1 /note="Met not found in wild type RapLRI"

PN W09950398-AZ.

PD 07-OCT-1999.

PF 26-MAR-1999; 99WO-US06641.

PR 27-MAR-1998; 98US-0079751.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Newton DL, Rybak SM;

DR WPI: 1999-610847/52.

DR N-PSDB: AAY08126.

PT New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases
 PS Claim 34; Page 57; 71pp; English.

CC The present sequence is a recombinant Rana pipiens ribonuclease (RapLRI)
 CC protein with Met at position 1. Carboxy terminal end of recombinant
 CC RapLRI has a covalently bound ligand binding moiety, which can be a IL2
 CC antibody directed against CD22 on cancerous B cells or human chorionic
 CC gonadotrophin (hCG) effective against Kaposi's sarcoma cells. Recombinant
 CC ribonucleases can be expressed in bacteria without an N-terminal
 CC methionine due to the presence of a signal peptide that is cleaved by
 CC bacteria. The soluble expression of ribonuclease allows the proteins to
 CC be fused in-frame with ligand binding moieties to form cytotoxic fusion
 CC proteins. They can be used for treatment of cancer and autoimmune
 CC diseases.

SO Sequence 105 AA;

Query Match 46.8%; Score 281.5; DB 20; Length 105;
 Best Local Similarity 49.5%; Pred. No. 1.4e-24;
 Matches 55; Conservative 15; Mismatches 32; Indels 9; Gaps 4;

OY 1 QNATFOQKHIIINT-PIICNTIMDNNTIYVGGCKRVNFTISSATVKAICTGVI-NMN 58
 DB 2 QDWLTFOQKHIIINT-PIICNTIMDNNTIYVGGCKRVNFTISSATVKAICTGVI-NMN 58
 OY 59 VLSSTRQALNTCTRTSTTPRCPPYSSRTETNYICVGCENQYPVHFGIGRC 109

DB 58 VLTTSERYLSDC---NWTSRPCKYKLLKSTNTFCVTCENQAPVHFGVGHG 105

RESULT 11

AAy28879
 ID AAY28879 standard; Protein: 127 AA.

AC AAY28879;
 DT 25-JAN-2000 (first entry)

DE Rana pipiens Clone 5a1b ribonuclease.

KW Rana pipiens ribonuclease Clone 5a1b; RapLRI; covalently bound; RNase;
 KM IL2 antibody; ligand binding moiety; CD22; cancerous B cell; onconase;
 KM Kaposi's sarcoma; human chorionic gonadotrophin; hCG; cancer;
 KM recombinant ribonuclease; cytotoxic fusion protein;
 KM autoimmune disease.

OS Rana pipiens.

FT Key Location/Qualifiers
 FT Peptide 1..23 /label=Signal-peptide
 FT Protein 24..127 /note="Putative"

FT /label= Rana_pipiens_Clone_5a1b_ribonuclease

PN W09950398-AZ.

PD 07-OCT-1999.

PF 26-MAR-1999; 99WO-US06641.

PR 27-MAR-1998; 98US-0079751.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Newton DL, Rybak SM;

DR WPI: 1999-610847/52.

DR N-PSDB: AAY08136.

PT New recombinant ribonucleases, used for killing target cells, e.g. for
 PT treating cancers, viral infections or autoimmune diseases
 PS Disclosure: Page 69; 71pp; English.

CC The present sequence is a Rana pipiens Clone 5a1b ribonuclease (RapLRI).
 CC It is encoded by Clone 5a1b cDNA obtained from Rana pipiens liver mRNA
 CC library. It exhibits differences with Onconase (RPM) at amino acid
 CC residues 11, 20, 85 and 103. Carboxy terminal end of RapLRI has a
 CC covalently bound ligand binding moiety, which can be a IL2 antibody
 CC directed against CD22 on cancerous B cells or human chorionic
 CC gonadotrophin (hCG) effective against Kaposi's sarcoma cells. Recombinant
 CC ribonucleases can be expressed in bacteria without an N-terminal
 CC methionine due to the presence of a signal peptide that is cleaved by
 CC bacteria. The soluble expression of ribonuclease allows the proteins to
 CC be fused in-frame with ligand binding moieties to form cytotoxic fusion
 CC proteins. They can be used for treatment of cancer and autoimmune
 CC diseases.

SO Sequence 127 AA;

Query Match 46.8%; Score 281.5; DB 20; Length 127;
 Best Local Similarity 49.5%; Pred. No. 1.8e-24;
 Matches 55; Conservative 15; Mismatches 32; Indels 9; Gaps 4;

OY 1 QNATFOQKHIIINT-PIICNTIMDNNTIYVGGCKRVNFTISSATVKAICTGVI-NMN 58
 DB 24 QDWLTFOQKHIIINT-PIICNTIMDNNTIYVGGCKRVNFTISSATVKAICTGVI-NMN 58
 OY 59 VLSSTRQALNTCTRTSTTPRCPPYSSRTETNYICVGCENQYPVHFGIGRC 109

Db 80 VLTISEFLSDC---NVTSRPCKYKLLKSTNTFCVTCENQAPVHFVGHC 127

RESULT 12

AAV28866 ID AAV28866 standard; Protein: 104 AA.

AAV28866; XX

25-JAN-2000 (first entry) XX

Recombinant RapLRI Met23leu amino acid sequence. XX

Recombinant Rana pipiens ribonuclease; RapLRI Met23leu; covalently bound; XX

KL2 antibody; ligand binding moiety; CD22; cancerous B cell; RNase; XX

Kaposi's sarcoma; human chorionic gonadotropin; hCG; signal peptide; XX

recombinant ribonuclease; cytotoxic fusion protein; cancer; frog; XX

autoimmune disease. XX

Rana pipiens. XX

Synthetic. OS

Key Location/Qualifiers XX

Misc-difference 23 /note= "Wild type Met replaced with leu" XX

MO9950398-A2. XX

07-OCT-1999. XX

26-MAR-1999; 99WO-US06641. XX

27-MAR-1998; 98US-0079751. XX

(USSH) US DEPT HEALTH & HUMAN SERVICES. XX

Newton DL, Rybak SM; XX

WPI: 1999-610847/52. XX

N-PSDB: AA208125. XX

New recombinant ribonucleases, used for killing target cells, e.g. for XX

treating cancers, viral infections or autoimmune diseases - XX

Claim 34; Page 56; 71pp; English. PS

The present sequence is a recombinant Rana pipiens ribonuclease (RapLRI) XX

protein with Met23leu. Carboxy terminal end of recombinant RapLRI has a XX

covalently bound ligand binding moiety, which can be a KL2 antibody XX

directed against CD22 on cancerous B cells or human chorionic XX

gonadotropin (hCG); effective against Kaposi's sarcoma cells. Recombinant XX

ribonucleases can be expressed in bacteria without an N-terminal XX

methionine due to the presence of a signal peptide that is cleaved by XX

bacteria. The soluble expression of ribonuclease allows the proteins to XX

be fused in-frame with ligand binding moieties to form cytotoxic fusion XX

proteins. They can be used for treatment of cancer and autoimmune XX

diseases. CC

Sequence 104 AA: SQ

Query Match 46.3%; Score 278.5; DB 20; Length 104;

Best Local Similarity 48.6%; Pred. No. 3e-24; Matches 54; Conservative 16; Mismatches 32; Indels 9; Gaps 4;

1 QNMTFPOKHLINT-PIICNTIMDNIIYVGQCKRVNFTIISATYKAICTGVI-NMN 58

1 QDMTLFQKHLINTRDVDCNNILSTNLF---HCKDKNTFTYSRPEPYKAICKGIASKN 56

59 VLTSTRPOLNCTRTSTIRPCPYSSRRETNYICVCKCNOYPVHFAGIGRC 109

57 VLTISEFLSDC---NVTSRPCKYKLLKSTNTFCVTCENQAPVHFVGHC 104

RESULT 13

AAV28869 ID AAV28869 standard; Protein: 105 AA.

AAV28869; XX

25-JAN-2000 (first entry) XX

Recombinant Met(-1) RapLRI Met23leu-(His)6 protein. XX

Recombinant Met(-1) Rana pipiens ribonuclease Met23leu-(His)6; RapLRI; XX

CD22; covalently bound; KL2 antibody; ligand binding moiety; RNase; XX

cancerous B cell; Kaposi's sarcoma; human chorionic gonadotropin; hCG; XX

signal peptide; recombinant ribonuclease; cytotoxic fusion protein; XX

cancer; frog; autoimmune disease. XX

Rana pipiens. OS

Synthetic. XX

Key Location/Qualifiers XX

Misc-difference 1 /note= "(His)6 histidine tag attached to N-terminal Met" XX

Misc-difference 1 /note= "Met not found in wild type RapLRI" XX

Misc-difference 24 /note= "Wild type Met replaced with leu" XX

MO9950398-A2. XX

07-OCT-1999. XX

26-MAR-1999; 99WO-US06641. XX

27-MAR-1998; 98US-0079751. XX

(USSH) US DEPT HEALTH & HUMAN SERVICES. XX

Newton DL, Rybak SM; XX

WPI: 1999-610847/52. XX

N-PSDB: AA208127. XX

New recombinant ribonucleases, used for killing target cells, e.g. for XX

treating cancers, viral infections or autoimmune diseases - XX

Claim 4; Page 59; 71pp; English. PS

The present sequence is a recombinant Rana pipiens ribonuclease protein XX

(RapLRI) with Met at position 1 attached to (His)6 tag and Met24leu. XX

Carboxy terminal end of recombinant RapLRI has a covalently bound ligand XX

binding moiety, which can be a KL2 antibody directed against CD22 on XX

cancerous B cells or human chorionic gonadotropin (hCG); effective XX

against Kaposi's sarcoma cells. Recombinant ribonucleases can be XX

expressed in bacteria without an N-terminal methionine due to the XX

presence of a signal peptide that is cleaved by bacteria. The soluble XX

expression of ribonuclease allows the proteins to be fused in-frame with XX

ligand binding moieties to form cytotoxic fusion proteins. They can be XX

used for treatment of cancer and autoimmune diseases. CC

Sequence 105 AA: SQ

Query Match 46.3%; Score 278.5; DB 20; Length 105;

Best Local Similarity 48.6%; Pred. No. 3.1e-24; Matches 54; Conservative 16; Mismatches 32; Indels 9; Gaps 4;

1 QNMTFPOKHLINT-PIICNTIMDNIIYVGQCKRVNFTIISATYKAICTGVI-NMN 58

2 QDMTLFQKHLINTRDVDCNNILSTNLF---HCKDKNTFTYSRPEPYKAICKGIASKN 57

59 VLTSTRPOLNCTRTSTIRPCPYSSRRETNYICVCKCNOYPVHFAGIGRC 109

58 VLTISEFLSDC---NVTSRPCKYKLLKSTNTFCVTCENQAPVHFVGHC 105

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XH XX Key Location/Qualifiers
FT Modified-site 1 /note= "this Gln is autocyclised to pyroglutamic acid"
XX PN US6175003-BI.
XX PD 16-JAN-2001.
XX PF 10-SEP-1999; 99US-0394268.
XX PR 10-SEP-1999; 99US-0394268.
XX PA (ALFA-) ALFACELL CORP.
XX PI Saxena SK;
XX DR WPI: 2001-167808/17.
XX PT New nucleic acids encoding a ribonuclease (Rnase), useful for the
XX precise targeting of Rnase to a predetermined cell receptor .
XX PS Claim 1; Columns 5-6; 7pp; English.

CC CC The present sequence represents a frog ribonuclease protein (rampirase)
CC (RNase). The specification describes a synthetic ribonuclease protein,
CC in which the addition of cysteine in the ribonuclease facilitates the
CC chemical linking of a targeting molecule by the single reactive
CC sulhydryl group. The specification also describes a method for the
CC production of rampirase using DNA technology instead of processing
CC biological material. The re-engineering of the protein molecule allows
CC easier attachment to a targeting molecule thereby making it possible for
CC the ribonuclease to be delivered to a particular cell receptor where it
CC might be most effective.
SQ Sequence 104 AA:
    Query Match          46.1%; Score 277.5; DB 22; Length 104;
      Best Local Similarity 49.5%; Pred. No. 4e-24;
        Matches 55; Conservative 15; Mismatches 32; Indels 9; Gaps 4
          1 QNMATFOQKHIINT-PIICNTIMDNIIYVGQCKRVMTFISSATVKAICTGYI-NMN 58
            I:I :| |:| | | | : | : | | : | | | | | | | | | | | |
          Dd 1 QDWLTFFOKHITNTRDVCNDINMSTLNF----HCKDKMTFITSRPEPVKAIKGIISKN 56
              1
          Oy 59 VLSTTROLNCTRTSITPRCPISSTRLETNYICVKCENQYPVHEAGIGRC 109
              ||::||:| | : | | | | | | | | | | | | | | | | | |
          Db 57 VLTISEFLSDC---NWTSRPCKYKLKKSTNKFCVTCENOAPVHFVGVCSC 104
              ||::||:| | : | | | | | | | | | | | | | | | | | |

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